

GROWTH STIMULATING COMPOSITION FOR PLANTS

The present invention refers to a product with the capacity to increase the intra-cellular levels of cyclic AMP as stimulants of plant growth and 5 development in normal conditions and in conditions of stress (osmotic, hydric, thermic, mechanical or pathogenic attack). It can be applied to any species of plant and at any point in its life cycle.

Although the intervention of cyclic AMP (3'-5' - 10 adenosine cyclic monophosphate) is well-known as an intra-cellular regulator of a great number of biochemical processes in animals (Khan y Lands, 1973. Prostaglandins and cyclic AMP. Academic Press), the role that this compound may represent in the regulation 15 of biochemical processes in plant cells is far from being well-known and even its existence in this medium is disputed (Newton, R., and col., Cyclic nucleotides in higher plants: the enduring paradox. New Phytol., 143, 427-455., 1999).

20 The effect of the intra-cellular levels of cyclic AMP on the growth of plants in stress conditions and normal conditions has been investigated. The stress states include osmotic or saline stress, hydric stress, thermic stress, mechanical stress and stress caused by 25 the attack of exterior agents (pathogenic agents like, for example, fungi, bacterias, insects, viruses, etc).

At this present time, some patent documents are known which describe the use of nucleotides to improve germination and primary development of seedlings.

30 In this way, in document US 4,209,316 (McDaniel and Brooks; 1978), it is not specified if they are cyclic nucleotides and furthermore the use of stimulants is not contemplated in the intra-cellular content of cyclic AMP but in the nucleotides 35 themselves. Moreover, neither do they refer to adult

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plants in production but to seedlings in their primary states.

5 Document US 4,050,919 (Motomura, Y and Ishiyama, J. 1975) refers to the direct application of cyclic AMP and not products which act on the intra-cellular content of cyclic AMP.

10 The same applies to other known patent documents, like for example document JP63279722A (Tanimoto Kiyoshi, Takahashi Shigeru; 1987) where it describes the application of cyclic AMP in culture media for the development of seedlings *in vitro*. Nevertheless, the application of cyclic AMP directly presents major problems due to the reduced capacity of this molecule 15 to penetrate through the biological membranes (Khan and Lands, 1973. Prostaglandins and cyclic AMP. Academic Press), making its effectiveness (in animal models too) very low.

20 One objective of the present invention is to have a compound to stimulate the growth of plants which can be used in any type of plant and at any point in its life cycle.

25 Another objective of the present invention is to have a compound to stimulate plant growth which is effective both in normal conditions and in stress situations (osmotic, hydric, thermic, mechanical or pathogenic attack).

30 The invention basically consists of using a precursor compound of **cyclic AMP**, or an inhibitor compound of the enzymes of the **phosphodiesterases** family, or a stimulator compound of the enzymes of the **Adenyl- Cyclase**, or an agonist compound of the  $\beta$ -adrenergic receptors to which arachidonic acid or a prostaglandin is added.

35 The advantages which the product object of the invention presents with respect to the state of the

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technique lie in:

- Its composition, since compounds are used (different from cyclic AMP) with the capacity to increase the intra-cellular content of cyclic AMP in 5 different species of plants, thus stimulating its plant development. None of the products specified have been previously described for this use in plants.

- Its use, since the product object of the invention is prepared for the treatment of adult plants in 10 production. The known patent documents only refer to culture media in relation to the production of plants in vitro, seed germination or primary development of seedlings.

- The conditions of use, since the product object of 15 the invention is active on adult plants both under normal conditions, **and under stress conditions** (saline, osmotic, hydric, mechanical or caused by pathogen agents).

A preferred embodiment of the composition object of 20 the invention comprises one or several of the following components;

A precursor compound of cyclic AMP for its transformation to the latter in the interior of the cells, such as the by-product Dibutiril-cyclic AMP.

25 The optimum dosage interval is between 0.1 and 2 ppm, via radicular or foliar.

A compound with the capacity of inhibiting the activity of the enzymes of the **phosphodiesterases** family, such as theophylline, theobromine or caffeine.

30 The optimum dosage interval is between 0.5 and 10 ppm, via radicular or foliar.

A compound with the capacity to stimulate the activity of the enzymes of the **Adenyl-Cyclase**, such as forskolin. The optimum dosage interval is between 0.1

35 and 2 ppm, via radicular or foliar.

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5 An agonist compound of the  $\beta$ -adrenergic receptors such as isoproterenol, epinephrine (adrenaline) or norepinephrine (noradrenaline). The optimum dosage interval is between 1 and 100 ppm, via radicular or foliar.

A chosen compound between arachidonic acid or a prostaglandin. The optimum dosage interval is between 0.25 and 100 ppm, via radicular or foliar.

10 As for the applicability of the composition object of the invention it can be brought into practice at great liberty.

The composition object of the invention can be applied in solid form, liquid form or suspension.

15 The composition object of the invention can be applied to any species of plant and at any point in its life cycle. The optimum time of application is:

- Via foliar or radicular in annual plants (cereal, horticultural,...): commence treatment on transplant or with 3-4 green leaves.

20 - In woody perennials commence treatment after budding, and before flowering.

The composition object of the invention can be applied by via radicular or foliar. The optimum dosage interval is between 5 and 10 litres or kilos per hectare. When applied in solution, the optimum dilution is 0.5%.

30 The composition object of the invention can be presented mixed with mineral fertilizers (such as amonic nitrate, monopotassium phosphate, etc.), with phyto regulators (such as cytoquinines, auxines, gibberelines, polyamines, n-ethanolamines, sugars ...) or any type of phytosanitary product (such as fungicides, herbicides, etc.). The proportion of each compound in the mixture can vary between:

35 1-25% for fertilizers

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0.5-5% phytoregulators

1-20% phytosanitaries.

The composition object of the invention can be formulated with tensoactive agents, such as Tween 80 5 etc., moistening agents etc. The proportions of the mixture of each compound can vary between:

0.1%-10% tensoactive.

1-10% moistening agents.

EXAMPLE

10 Composition for 1 kg of product:

1. 0.5 g of Forskolin

2. 1 g of caffeine

3. 0.2 g of 6-Bencilaminopurina

4. 200 g of dipotassium phosphate

15 5. 45 g of urea

6. 753.3 g of water

Order of making:

Continuous stirring and environment temperature (20-25°C)

20 1. (6)+(4)+(5). Stirring until completely dissolved.

2. 1.+(1)+(2)+(3).

Effect of different compounds on the development of vine plants affected by an attack of mildew 25 (plamopara viticola).

A random block assay was carried out with four repetitions per treatment including 10 vines per repetition.

The disease caused by the plamopara viticola 30 fungus was evaluated both in the expression of frequency (number of leaves affected) and of intensity (percentage of leaf surface affected).

The products applied were:

35 - Treatment A: A by-product of phosphorous acid + a contact fungicide (folpet) (4kg/hectare) (reference

treatment).

- Treatment B: A by-product of phosphorous acid (2.35 l/hectare).

5 - Treatment C: A by-product of phosphorous acid (3.25 l/hectare) + Patent product according to example of making (2 l/hectare).

- Control which received no treatment.

The results were the following

Degree of disease on leaves

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<u>Treatment</u>	<u>Frequency (%) (% reduction)</u>	<u>Intensity (%) (% reduction)</u>
Control	12.27 a (-)	53 a (-)
A	0.5 b (96)	4.8 b (91)
B	4.7 c (62)	25.8 c (51)
C	1.85 b (85)	11.8 b (78)

As can be observed, the application of the product object of the patent with the capacity to increase the intra-cellular levels of cyclic AMP (product C) strengthens the action of the product of synthesis (by-product of phosphorous acid) to 23% in relation to

15 frequency and to 27% in relation to intensity referring to the absolute value of the % of reduction.